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(54) **STABLE HYBRID**

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(57) **ABSTRACT**

A hybrid formed of a probe nucleic acid immobilized onto a substrate and a sample nucleic acid, in which the target strand of the sample nucleic acid is longer than the probe nucleic acid, so that at least a 5' end portion of the target strand extends upstream the target sequence; a 3' end of the target strand is the same as a 3' end of the target sequence or extends downstream the target sequence by at least one base from the 3' end of the target sequence; and provided that when the number of bases of the extending 5' portion is designated as L1, and the number of bases of the extending 3' portion of the target strand is designated as L2, a value of L1/L2 falls within the range of 0 to 1.5 both inclusive. This hybrid is very stable so that useful for detecting a small amount of nucleic acid specifically with high sensitivity.

**2 Claims, 4 Drawing Sheets**

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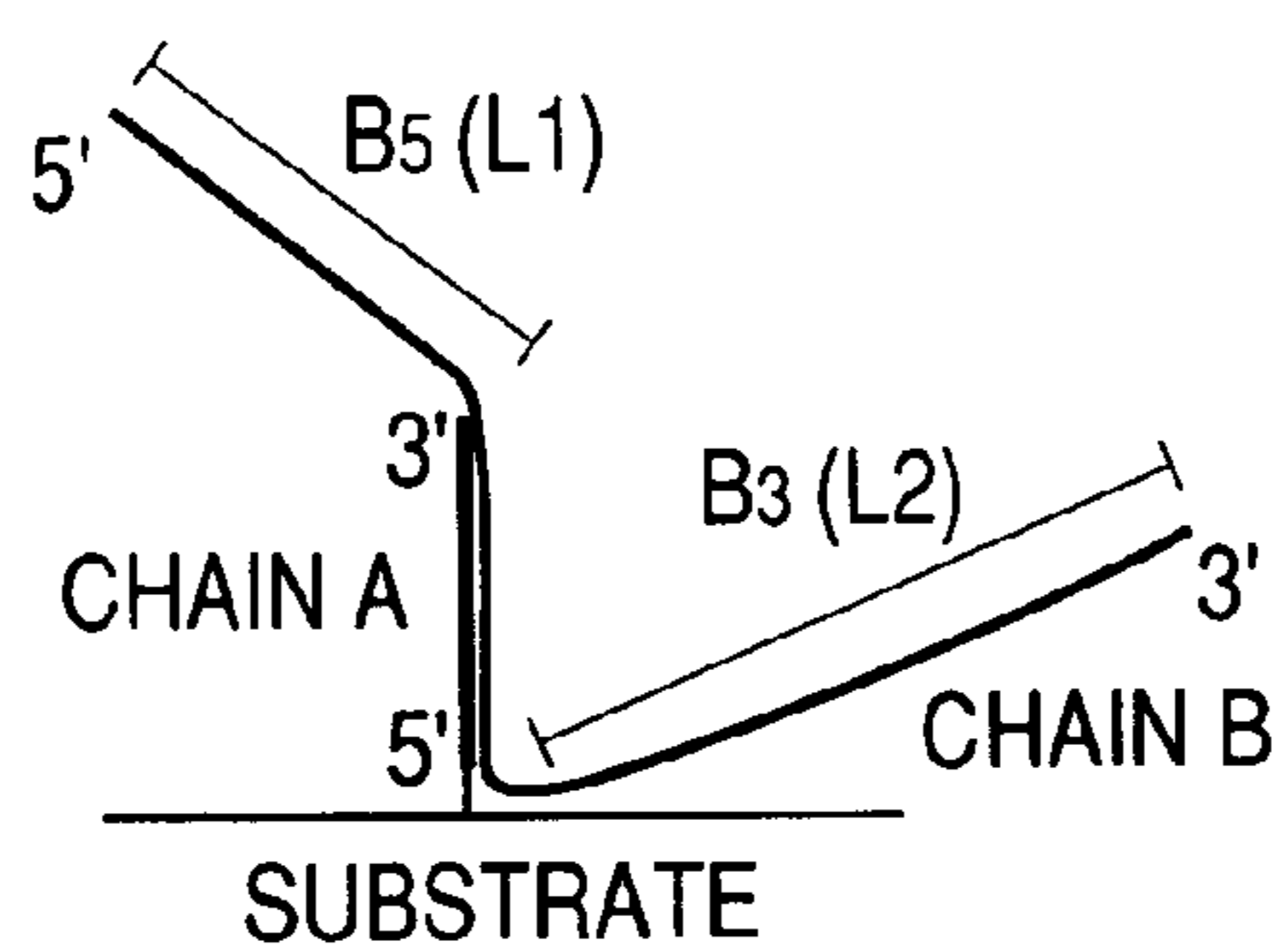
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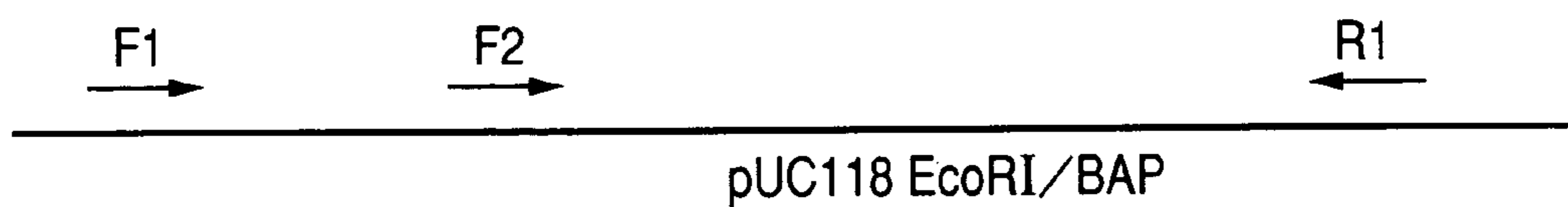
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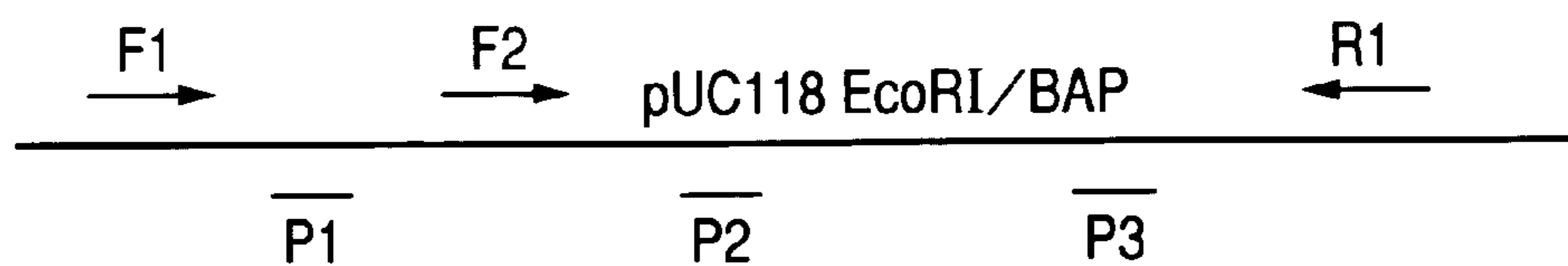
**FIG. 1**



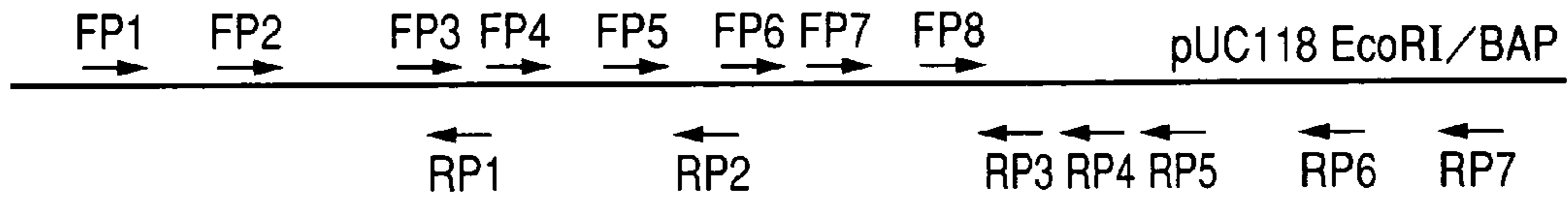
**FIG. 2**



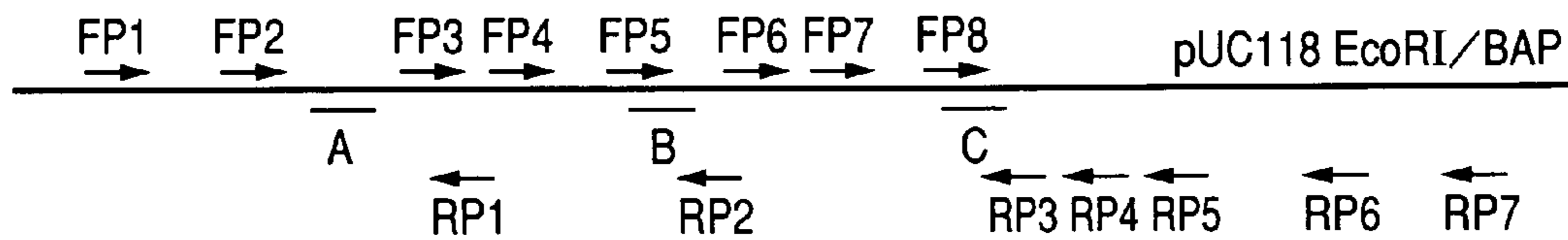
**FIG. 3**



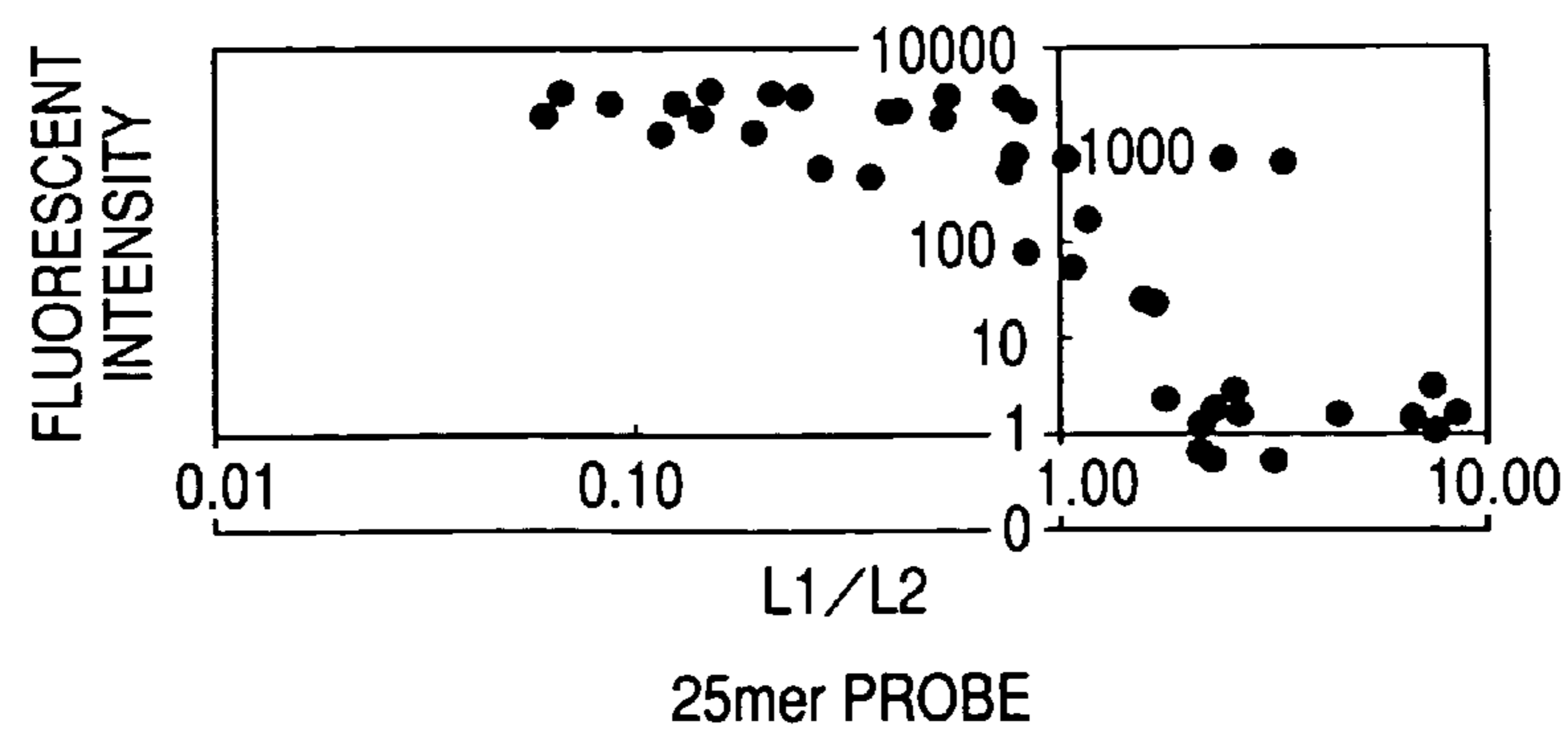
*FIG. 4*



*FIG. 5*

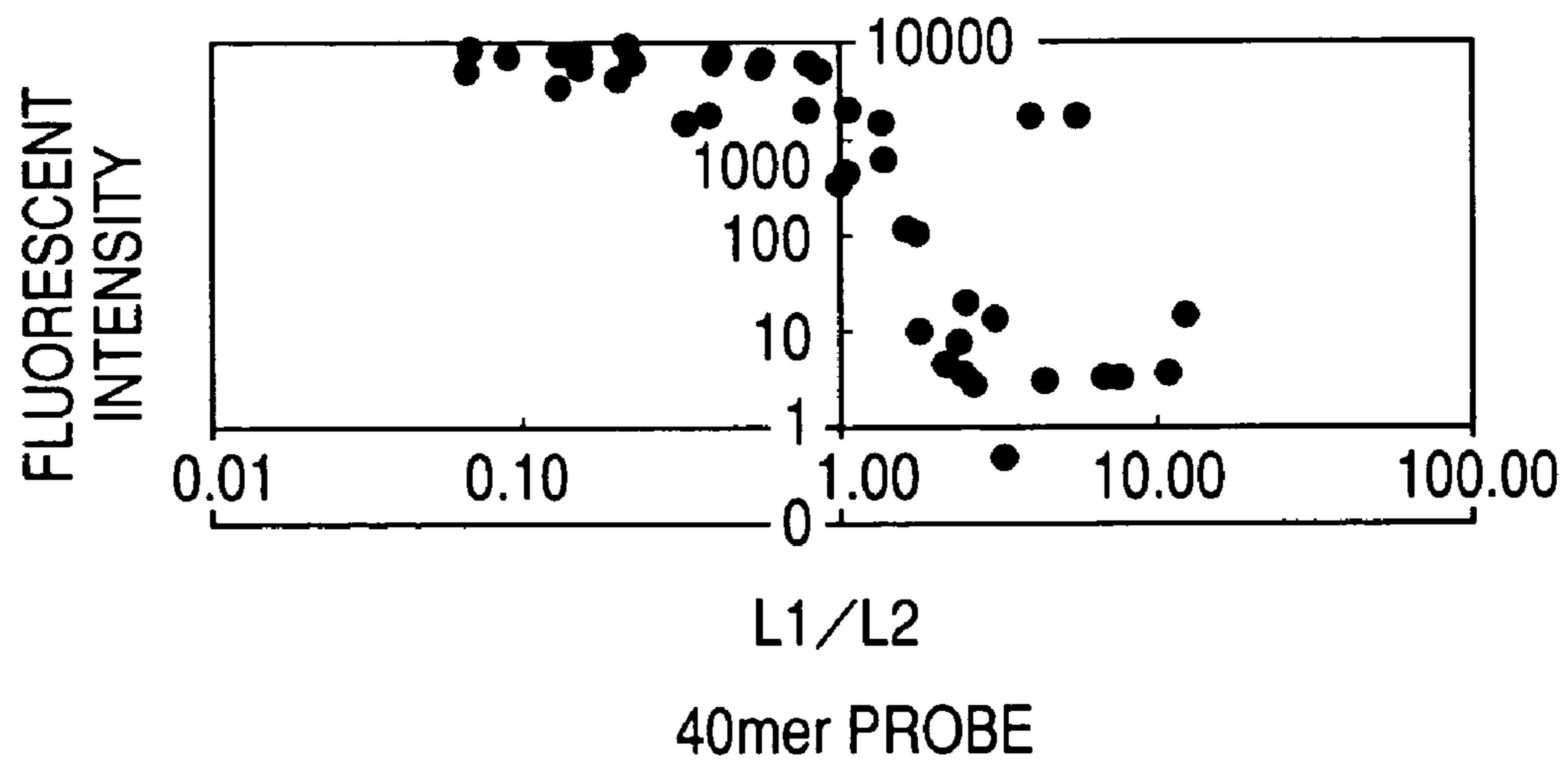


*FIG. 6*

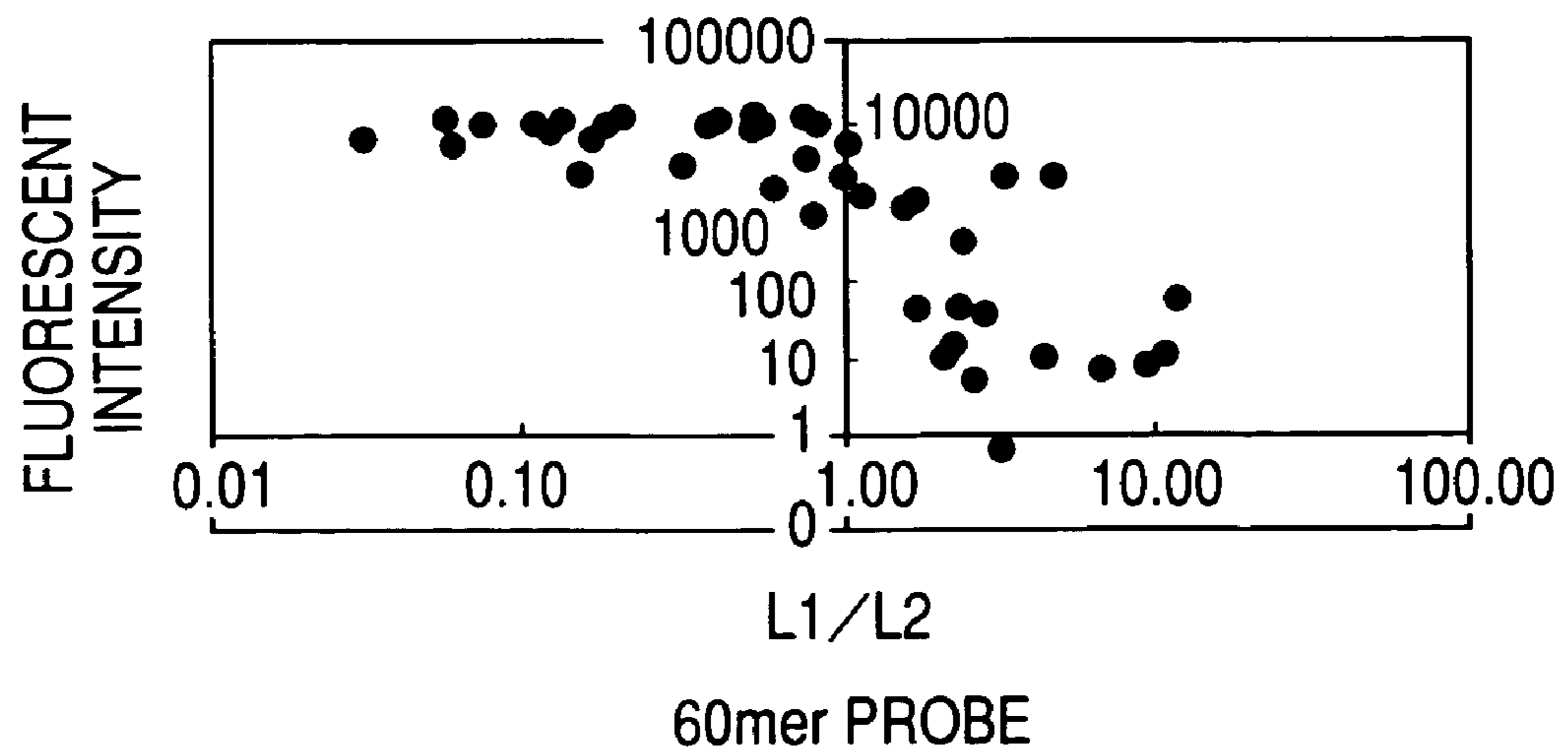




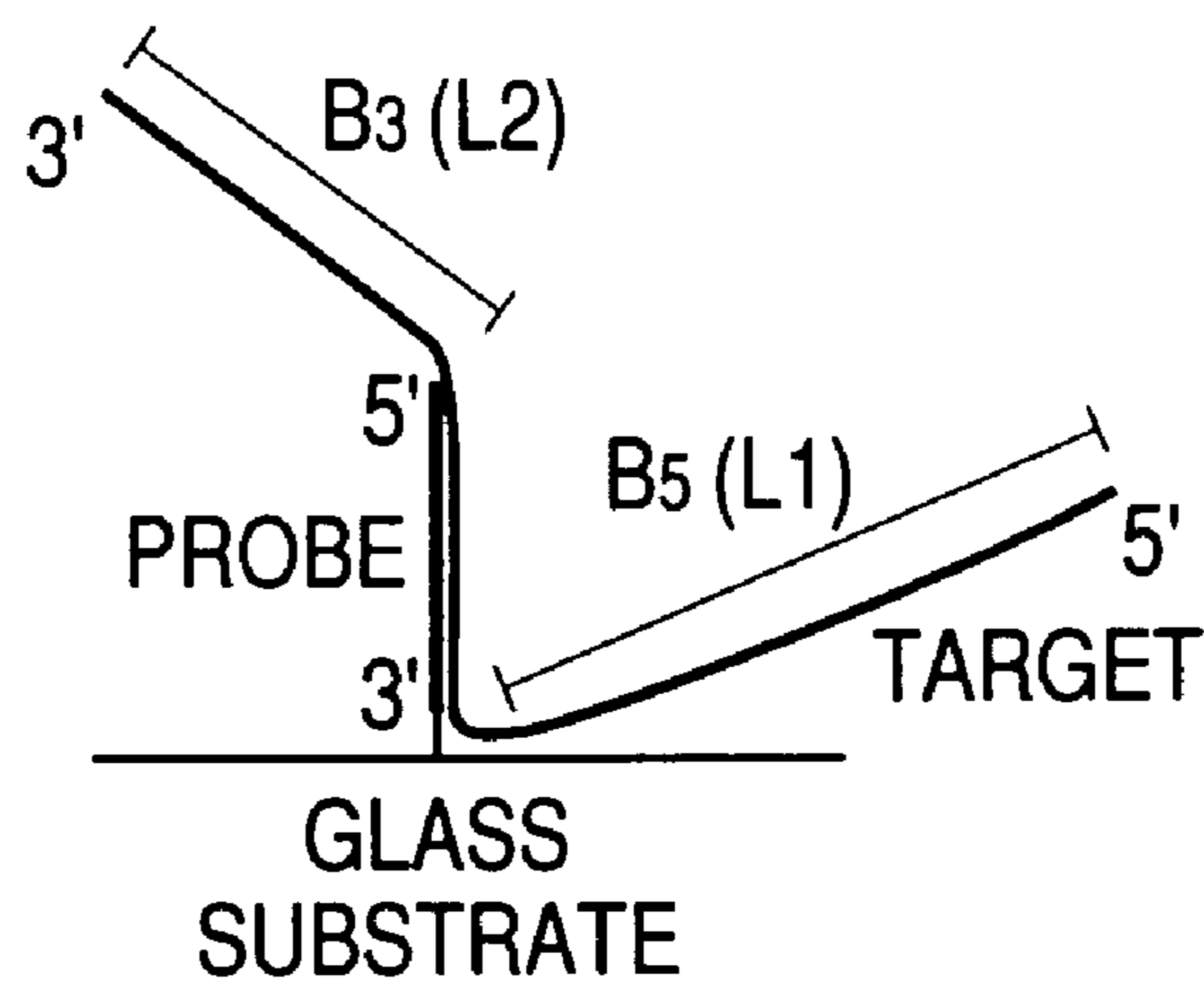
*FIG. 7*



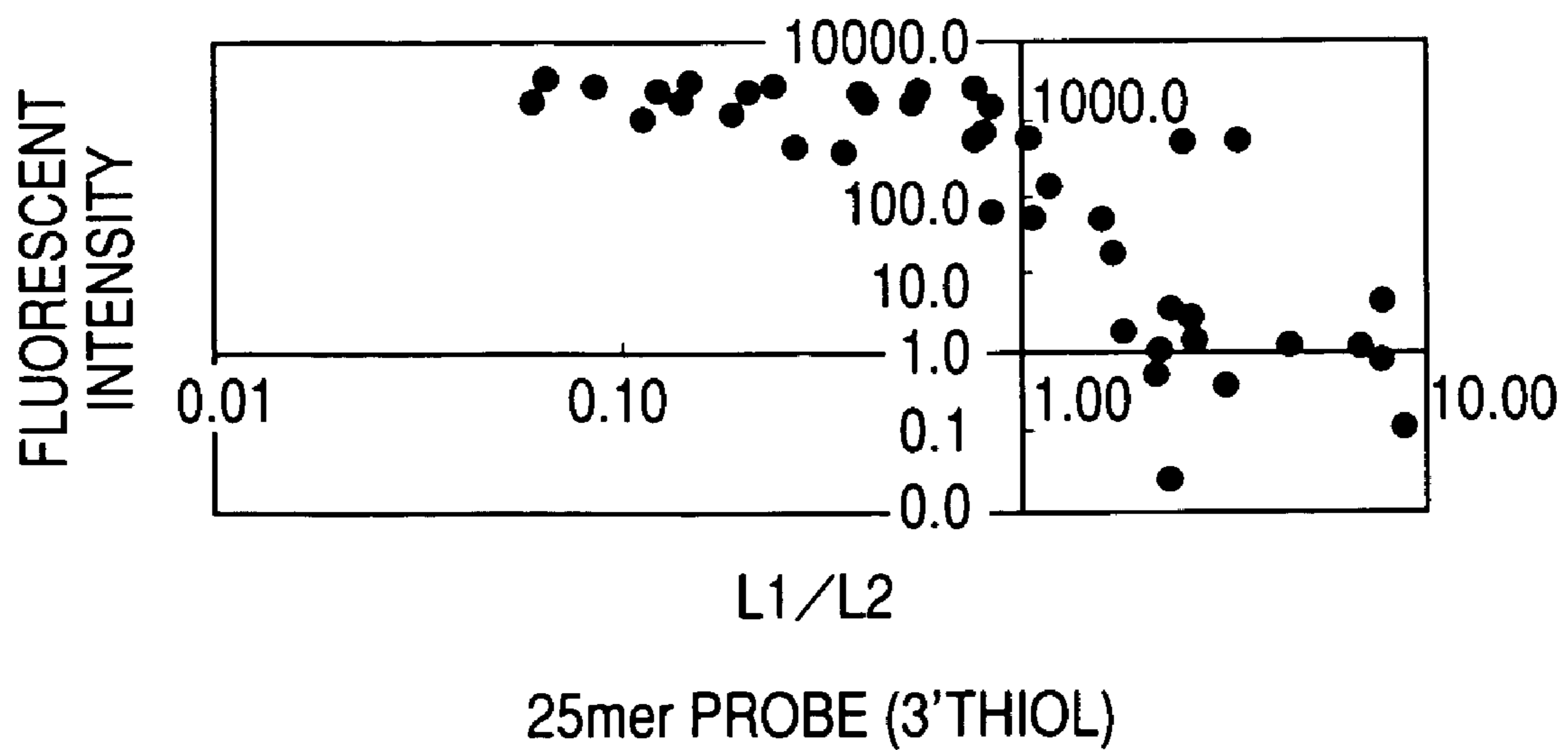
*FIG. 8*



**FIG. 9**



**FIG. 10**





## STABLE HYBRID

## BACKGROUND OF THE INVENTION

## 1. Field of the Invention

The present invention relates to formation of a stable hybrid between a sample nucleic acid and a probe immobilized on a substrate. The present invention also relates to a probe for use in forming such a hybrid, a PCR primer, a probe carrier and designing methods thereof.

## 2. Related Background Art

As is represented by the Human Genome Project, gene sequences of various types of organisms have been elucidated and the relationships between genes and the mechanism of life activity, diseases, constitutional predispositions and so forth are investigated. It is now understood that by determining the presence or absence of a certain gene or its (expression) amount, not only the feature or type of a disease but also the effective therapy thereof may be determined.

Various methods have been proposed for detecting the presence or absence of a specific gene in a test sample and determining the expression amount of the gene, among which, the following method is applicable to a wide variety of samples. This method comprises the steps of selecting (determining) a partial nucleotide sequence specific to a target gene, and detecting the presence or absence or the expression amount of the partial sequence, thereby determining the presence or absence of the gene or its expression amount. More specifically, according to this method, a nucleic acid (probe) of which sequence is complementary to the selected partial sequence is prepared, and the probe and a test sample are hybridized to detect whether a hybrid is formed or not by certain means, whereby the presence or absence of the gene in the sample is determined.

This method of detecting a specific nucleic acid by using hybridization can be used in both solid and liquid phases. For example, hybridization in a liquid phase is performed as follows: a probe labeled with a labeling agent of which spectroscopic property changes when the probe becomes double stranded; then a test sample is added to a solution of the probe; and spectroscopic change is measured. In this way, the presence or absence of a specific nucleic acid can be determined.

On the other hand, hybridization on a solid is typically performed as follows: a probe is immobilized onto a substrate; then, a sample labeled with a detectable labeling agent is applied to the substrate; and a signal emitted from the labeling agent on the substrate is measured. Typical examples are microarrays where probes are immobilized on a flat glass or metal substrate, and beads that are fine particles having immobilized probes thereon. Solid-phase hybridization is preferably used for the following reasons: B/F separation (separation of bound form and free form) is easy; detection can be performed in a minute area and with high sensitivity; if a plurality of types of probes are discretely arranged, simultaneous detection of a plurality of items can be made; and solid substrates are easy to handle or use.

For example, U.S. Pat. No. 6,410,229 to Affymetrix Inc. (USA) discloses a method for detection and quantification of a specific nucleic acid in a sample by hybridizing a labeled nucleic acid with an oligo DNA synthesized on a flat substrate, and detecting fluorescence emitted from the hybrid. On the other hand, Japanese Patent Publication No. 2001-128683 (Fuji Photo Film Co., Ltd.) discloses detection of 22 mer ssDNA by using a DNA array prepared using a substrate to which amino groups had been introduced beforehand.

## SUMMARY OF THE INVENTION

As described above, the detection method using solid-phase hybridization has an advantage over other detection

methods in sensitivity. However, the sensitivity of conventional solid-phase hybridization detection methods is not enough to satisfy needs for detecting a very small amount of nucleic acid. To detect a hybrid with higher sensitivity and higher specificity, improvement in the solid-phase hybridization detection method is demanded.

To increase detection accuracy, improvement has been made from various aspects. One method for improving the detection performance is to increase binding between the probe and the target by increasing the  $T_m$  of the probe-target. Alternatively, labeling rate of the target with a labeling agent is increased, a labeling agent emitting a strong signal is used, or the labeling agent itself is labeled (multi-labeling), to increase sensitivity.

However, the method of increasing binding of a probe to a target has a problem in that nonspecific adsorption and binding take place, greatly reducing specificity of target binding. Therefore, the effect of this method is limited. On the other hand, the method of improving labeling has a problem in that the S/N ratio and quantitateness may decrease. The effect of this method is also limited.

Another method for improving hybridization efficiency is to stabilize the hybrid formed between a probe and a target. To stabilize the hybrid, it is necessary to optimize the hybridization conditions, i.e., salt concentration and concentration of the denaturing agent. However, this method also has a problem in that specificity and S/N ratio may decrease.

In these circumstances, it has been strongly desired to develop a method of forming a stable hybrid, a probe designing method and a sample preparation method in order to detect a nucleic acid more efficiently.

The present inventors have intensively studied the aforementioned problems, and found out that a probe and a sample nucleic acid can hybridize to form a stable hybrid if a certain relationship between the probe immobilized on a substrate and the sample nucleic acid is satisfied.

The present invention provides a hybrid formed between a probe nucleic acid and a sample nucleic acid, wherein the probe nucleic acid and the sample nucleic acid are hybridized between a target sequence present in a target strand of the sample nucleic acid and a complementary probe sequence present in the probe nucleic acid, wherein the target strand of the sample nucleic acid is longer than the probe nucleic acid, and when the number of bases of a 5' portion of the target strand not being the target sequence is designated as L1, and the number of bases of a 3' portion of the target strand not being the target sequence is designated as L2, L1 and L2 satisfy the following relationship:  $0 < L1/L2 < 1.5$ .

The present invention also provides a probe designed so as to form the above-described hybrid with a target strand having a target sequence.

The present invention also provides a probe set comprising not less than two probes each being a probe described above.

The present invention also provides a probe carrier having probes for detecting a target nucleic acid immobilized, wherein not less than 50% of all probes is occupied by the above probe set. The present invention also provides a method of designing a probe so as to form the above-described hybrid with a target strand having a target sequence.

The present invention also provides a PCR primer for preparing a sample nucleic acid capable of forming a hybrid that is described above with a probe that is described above.

The present invention also provides a PCR primer for preparing a sample nucleic acid capable of forming an above-described hybrid with an above-described probe.

The present invention also provides a method of detecting a sample nucleic acid by detecting a hybrid obtained by



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reacting a sample nucleic acid with a probe nucleic acid immobilized in a substrate, the sample nucleic acid being longer than the probe nucleic acid, wherein the sequence of the probe nucleic acid is designed so as to form a hybrid satisfying features (1) and (2):

(1) the hybrid is formed between a target sequence present in a target strand of the sample nucleic acid and a complementary probe sequence present in the probe nucleic acid; and  
 (2) the target strand of the sample nucleic acid is longer than the probe nucleic acid, and when the number of bases of a 5' portion of the target strand not being the target sequence is designated as L1, and the number of bases of a 3' portion of the target strand not being the target sequence is designated as L2, L1 and L2 satisfy the following relationship:

$$0 \leq L1/L2 \leq 1.5$$

When the sample nucleic acid is a product amplified by PCR using a sample nucleic acid, it is desirable that the PCR primer herein and the probe are designed such that they can form a hybrid satisfying the features (1) and (2).

The present invention also provides a kit for detecting a sample nucleic acid comprising

- a PCR primer for amplifying a sample nucleic acid having a target strand having a target sequence, from the test-sample nucleic acid, and
- a probe nucleic acid immobilized onto a substrate and prepared for determining the presence and absence of the target sequence in an amplified product which is amplified using the PCR primer and the test-sample nucleic acid as a template;

wherein the sequences of the primer and the probe nucleic acid are designed so as to form a hybrid satisfying features (1) and (2):

(1) the hybrid is formed between a target sequence present in a target strand of the sample nucleic acid and a complementary probe sequence present in the probe nucleic acid; and  
 (2) the target strand of the sample nucleic acid is longer than the probe nucleic acid, and when the number of bases of a 5' portion of the target strand not being the target sequence is designated as L1, and the number of bases of a 3' portion of the target strand not being the target sequence is designated as L2, L1 and L2 satisfy the following relationship:

$$0 \leq L1/L2 \leq 1.5.$$

A hybrid according to the present invention may be formed of any combination of a probe nucleic acid and a sample nucleic acid as long as the probe nucleic acid is immobilized onto a substrate. A preferable example of the present invention is a hybrid formed by immobilizing an oligonucleotide or DNA such as cDNA on a flat substrate as a probe, e.g., DNA microarray, and hybridizing a sample nucleic acid labeled with a detectable marker with the probe.

A probe may be immobilized onto a substrate by various methods, such as covalent bond, ionic bond and adsorption. A hybrid of the present invention can be formed no matter which immobilization method is employed.

Of these immobilization methods, covalent bond is preferable in view of forming a stable hybrid because it relatively not influenced by various conditional changes associated with hybridization such as heating and high salt concentration. As covalent bond, any type of bonding is applicable without limitation as long as bonding is made between a functional group capable of being introduced into a substrate and a functional group capable of being introduced into a nucleic acid probe. When a synthetic oligonucleotide is used as a probe, a thiol group, amino group, or a derivative thereof is frequently used as a functional group for use in modifica-

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tion since they can easily introduced. Of various methods, a method of introducing the functional group as mentioned above into the 5' end and the 3' end of an oligonucleotide is convenient in view of a synthetic process. A probe, which is immobilized on a substrate via such a functional group introduced to one end of an oligonucleotide, can be preferable used to form a hybrid according to the present invention.

In addition to the functional groups introduced to the 5' end or the 3' end, a functional group introduced into the middle portion of an oligonucleotide sequence, or a functional group of an artificial nucleotide derivative introduced into a sample nucleic acid during synthesis may be used for immobilization.

Since a nucleic acid has negative charge as a whole owing to the phosphate ester bonds in the skeleton of a nucleic acid, it can be adsorbed to a positively charged coating material introduced onto a substrate or to the substrate itself.

A probe to be immobilized onto a substrate may differ in chain length. The specificity and binding force of a probe varies depending upon chain length of the probe. Oligonucleotides of 15 to 30 mers, 31 to 50 mers and 51 to 80 mers are frequently used as a probe to be immobilized onto a substrate. They are also suitable for a hybrid according to the present invention.

Although nucleic acid such as oligonucleotide is generally used as a probe, a peptide nucleic acid (PNA), to which strong hybridization can be expected, may be used. In that case, more stable hybrids can be obtained, by satisfying the relationship between L1 and L2 according to the present invention.

More specifically, as long as the value of L1/L2 falls within the range of 0 to 1.5, a stable hybrid according to the present invention can be formed. A more stable hybrid can be formed in particular, when the value of L1/L2 falls within the range of 0 to 1.

The sample nucleic acid herein means nucleic acid having a target sequence that hybridizes with a probe and subjected to a reaction with the probe. It may be the test sample itself, or a nucleic acid fragment taken out from the sample by any means. The target sequence is not particularly limited, as long as it can specify a sequence that the test sample is presumed to contain. Depending upon the type of sample nucleic acid, the target sequence may be a part or whole of the sample nucleic acid to be detected in a test sample, or a complementary sequence thereof. The sample nucleic acid has a target strand containing the target sequence (complementary to a probe sequence) to be hybridized with a probe. Any type of nucleic acid, for example, double stranded DNA, single stranded DNA, and RNA, may be applied to the present invention, as long as it can hybridize with a probe immobilized onto a substrate under appropriate conditions. When the sample nucleic acid is a single strand DNA, the DNA itself is a target strand.

A method of preparing a sample nucleic acid is not particularly limited and a PCR amplification method and reverse transcription method of RNA are often used. A PCR product and reverse transcription product obtained by these methods may be preferably used in order to form a hybrid of the present invention.

When solid-phase hybridization is used for detecting a specific base sequence, as is in the case of a DNA microarray, sample nucleic acid is usually labeled by binding a labeling agent. A sample nucleic acid in which a labeling agent is incorporated is also preferably applied to the present invention for forming a hybrid.

As a labeling agent, a fluorescent substance is generally used since it is easy to handle and detectable with high sen-



sitivity. Also, a radioisotope can be preferably used as a marker since it has excellent sensitivity. Although there are great many choices for fluorescent substances, any one of fluorescent substances may be used. Of them, fluorescent dyes such as Cy3 and Cy5 are widely used and can be preferably used in a hybrid according to the present invention.

The length of a sample nucleic acid is not particularly limited and any length of a nucleic acid may be used to form a hybrid. Of them, a sample nucleic acid having a chain length of not less than 500 bp is preferable since a stable hybrid can be formed. A more stable hybrid may be formed if the chain length of a sample nucleic acid falls within the range of 500 bp to 2500 bp.

A designed probe is also provided in the present invention in order to hybridize with a sample nucleic acid to form the hybrid mentioned above. More specifically, the present invention provides a probe that is designed by setting a probe-capturing region in a sample nucleic acid at a position at which a hybrid according to the present invention can be formed, thereby forming a hybrid satisfying the positional relationship between a solid-phase probe and a sample nucleic acid.

The present invention also provides a probe set having not less than two different types of probes. To provide such probes, it is necessary to design the sequence of a probe. Therefore, a method of designing a probe is also provided.

In order to use a probe efficiently for detecting a target nucleic acid (target sequence), the probe is immobilized onto a solid carrier to facilitate B/F separation. Therefore, the present invention provides a probe carrier having at least one probe is immobilized on a solid carrier. As an example of such a carrier, a DNA microarray may be mentioned. Such a DNA microarray is one of preferable examples for a probe immobilized carrier according to the present invention.

As a sample nucleic acid, a PCR product may be often used as mentioned above. To prepare a PCR product, a PCR primer is used. Therefore, the present invention provides a PCR primer designed so as to satisfy the conditions of L1 and L2.

According to the present invention, it is possible to form a stable hybrid between a probe immobilized onto a substrate and a sample nucleic acid, and obtain the conditions in which the hybrid is formed. Furthermore, it is possible to provide a probe and a primer satisfying the conditions of the present invention, and further provide a method of designing such a probe and a primer, and a probe carrier on which such a hybrid is formed. According to the present invention, a nucleic acid can be detected efficiently with high sensitivity by a method of detecting a nucleic acid by use of hybridization performed on a substrate.

Other features and advantages of the present invention will be apparent from the following description taken in conjunction with the accompanying drawings, in which the same reference characters designate the same or similar parts throughout the figures thereof.

#### BRIEF DESCRIPTION OF THE DRAWINGS

The accompanying drawings, which are incorporated in and constitute a part of the specification, illustrate embodiments of the invention and, together with the description, serve to explain the principles of the invention.

FIG. 1 shows an illustration of a hybrid between a probe (Strand A) immobilized on a substrate and a sample nucleic acid (Strand B) in which the length of 5' portion upstream the hybridized region of the sample nucleic acid is designated as  $B_5$ , and the length of 3' portion downstream the hybridized region as  $B_3$ ;

FIG. 2 shows the positions of three primers to pUC118 EcoRI/BAP in which the arrow of each primer is directed from the 5' to 3';

FIG. 3 shows three primers and three probes to pUC118 EcoRI/BAP;

FIG. 4 shows the positional relationship between 15 types of primers including forward primers F1 to F8 and reverse primers R1 to R7 to pUC118 EcoRI/BAP, in which the arrow of each primer is directed from the 5' to 3';

FIG. 5 shows the positional relationship between each of the primers and three probes to pUC118 EcoRI/BAP;

FIG. 6 is a graph plotting fluorescent intensity to L1/L2 value in the case of a 25-mer probe;

FIG. 7 is a graph plotting fluorescent intensity to L1/L2 value in the case of a 40-mer probe;

FIG. 8 is a graph plotting fluorescent intensity to L1/L2 value in the case of a 60-mer probe;

FIG. 9 shows an illustration of a hybrid between a probe immobilized at the 3' end and a target strand and the positional relationship between L1 and L2; and

FIG. 10 is a graph plotting fluorescent intensity to L1/L2 value in a 25-mer probe immobilized at the 3' end.

#### DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

Preferred embodiments of the present invention will now be described in detail in accordance with the accompanying drawings.

A hybrid according to the present invention will be explained in more detail.

A hybrid provided by the present invention is formed between a probe and a sample nucleic acid having a complementary sequence to the probe in part and satisfies the following relationship:

$$0 \leq L1/L2 \leq 1.5$$

wherein the chain length (base number) of 5' portion upstream the target sequence of the sample nucleic acid is designated as L1, and that of 3' portion downstream the target sequence as L2.

FIG. 1 shows a schematic view of an example hybrid of the present invention between a single stranded probe (Chain A) immobilized on a substrate at the 5' end and a single stranded nucleic acid (Chain B) having a complementary sequence to the Chain A. In this figure, the chain length of 5' portion upstream the target sequence of the chain B designated as  $B_5(L1)$ , and that of 3' portion downstream the target sequence designated as  $B_3(L2)$  satisfy the following relationship:

$$B_5 \leq B_3.$$

If the value of L1/L2 ranges from 0 to 1.5, a stable hybrid of the present invention can be formed. To obtain more stable hybrid, it is desirable to set an L1/L2 value within the range of from 0 to 1.

It is possible to form a hybrid of the present invention as long as a probe is immobilized on a substrate or a solid-phase surface no matter how the probe is immobilized. A DNA microarray having a nucleic acid immobilized on a glass substrate is a representative example. Various types of DNA microarrays are applicable to forming a hybrid of the present invention. A material for a substrate includes a resin, metal, metal film, and fiber other than glass. As to form of a substrate, any form may be used including fine particles, a tip of an optical fiber, porous material, and fiber.

Any binding means such as adsorption and chemical bond may be used to immobilize a probe for hybrid formation. For



example, ionic bond is mentioned in the section "SUMMARY OF THE INVENTION". If a glass substrate or a resin surface is coated to have amino groups in advance, unmodified nucleic acid can be immobilized by ionic bond on the substrate or resin surface by only applying it thereto.

Alternatively, if a modified oligonucleotide having a functional group, such as a thiol group or amino group, introduced to the 5' end, 3' end or the middle portion of a nucleic acid, the nucleic acid can be immobilized on a substrate through a reaction of the introduced functional group. For example, in case of amino group, a succinimide group, which efficiently reacts with the amino group, is introduced to the solid-phase surface in advance, and an oligonucleotide is modified with the amino group at the 5' end, 3' end, etc., whereby covalent bond is easily formed between the oligonucleotide and the solid-phase surface simply by applying the oligonucleotide thereto. The oligonucleotide thus bound on the solid-phase surface can be suitably used as a probe for forming a hybrid of the present invention.

When a thiol group is used, similarly maleimide group is introduced to the substrate, an oligonucleotide having thiol group can be easily covalently bonded to the substrate. The oligonucleotide thus bound on the substrate can be suitably used as a probe for forming a hybrid of the present invention.

Besides DNA, PNA can be used as a probe. Also cDNA may be used as a probe. However, in order to form a hybrid more accurately, a chemically synthesized oligonucleotide is frequently used as a probe. Such a probe desirably has a base chain length of 15 to 30 mers, 31 to 50 mers, or 51 to 80 mers, to be used appropriately according to their characteristics.

A sample nucleic acid for use in forming a hybrid of the present invention is not particularly limited and may be used as long as it is longer than a probe (immobilized on a substrate) in terms of chain length.

A hybrid of the present invention is suitable for detecting a nucleic acid sequence on a DNA microarray or beads. In this case, as a sample nucleic acid, a PCR amplification product, a transcription product, reverse transcription product, and so forth are usually used. These nucleic acids are preferably used to form a hybrid of the present invention. To explain more specifically, a sample nucleic acid may be either double stranded or single stranded and either DNA or RNA to form a hybrid of the present invention.

Furthermore, when formation of a hybrid is detected by a spectroscopic means, a sample nucleic acid must be labeled with a labeling agent such as a fluorescent substance. A hybrid of the present invention can be formed even if a nucleic acid labeled with such a labeling agent is used. Of the fluorescent labeling agents, Cy dyes including Cy3 and Cy5 (manufactured by Amersham Biosciences) are frequently used. A sample nucleic acid including a fluorescent substance is applicable to form a hybrid of the present invention without any problem.

As a labeling agent having the same detection sensitivity as a fluorescent substance, <sup>32</sup>P radioisotope is often used. In this case, a hybrid of the present invention can be sufficiently formed.

A hybrid of the present invention can be formed regardless of the chain length of the sample nucleic acid. When the length of a sample nucleic acid is 500 bp or more, more stable hybrid can be formed. When the length of a sample nucleic acid is from 500 bp to 2500 bp, further more stable hybrid can be formed.

When a sample nucleic acid is, for example, a double stranded DNA, a sample nucleic acid may be double stranded with its complementary strand except for the hybridized region. Even in this case, a hybrid of the present invention can

be formed. More specifically, if there are portions satisfying the aforementioned relationship of L1 and L2 in the hybrid formed between the immobilized probe on a substrate and the sample nucleic acid, the hybrid is a highly stable hybrid according to the present invention.

In the case where the double stranded portion formed between a probe and a sample nucleic acid is shorter than the length of the probe, that is, the probe containing a mismatch region is hybridizing with a sample nucleic acid, a stable hybrid can be formed. In this case, provided that the length of a sample nucleic acid between the 5' end to the end of a double stranded portion is represented by L1 and the length of the sample nucleic acid between the 3' end to the other end of the double stranded portion is represented by L2, if L1 and L2 satisfy the aforementioned relationship between L1 and L2, a hybrid of the present invention can be formed.

Usually, probes for detecting the presence or absence of a target nucleic acid based on a specific hybrid formation between the probe immobilized on a substrate and a target are designed so as to detect the sample nucleic acid specifically with high sensitivity. Since most of such probe designing methods consider only the base sequence, effective probes for liquid-phase hybridization can be designed, but design of probes for solid-phase hybridization is not always satisfactory. Therefore, the hybrid formation conditions of the present invention give an indication to the stability of a hybrid on a substrate. Accordingly, the present invention also provides a method of designing a probe on a solid substrate that gives stable hybrid of the present invention.

The present invention further provides a probe designed in accordance with this probe designing method. Moreover, the present invention provides a set of probes containing a plurality of (two or more) probes.

A DNA microarray is an example of a probe carrier in which a plurality of different sequence of probes are immobilized on a solid carrier. When a probe of the present invention is used as probes constituting such a probe carrier, not less than 50% of all probes on the carrier is preferably occupied by probes according to the present invention, and not less than 70% is more preferable. Such a probe carrier having a plurality of probes (probe set) immobilized on a carrier is also provided by the present invention.

The probes satisfying the conditions of the present invention are preferably present in a ratio of 50% or more, and more preferably, in a ratio of 75% or more to the entire number of probes of a probe carrier.

When a PCR product is used as a sample nucleic acid, the relative position of the region corresponding to the probe (probe region) in the sample nucleic acid may vary depending upon the design of primers for PCR amplification. It is ideal that the probe region of a sample nucleic acid is designed so as to satisfy the aforementioned conditions (represented by the equation). Sometimes a probe region does not meet the conditions, for example, in view of specificity. In such a case, first a probe region is designed and then a primer for a sample nucleic acid may be designed such that the probe region satisfies the condition of the present invention. Therefore, the present invention also provides a primer for amplifying a sample nucleic acid having a predetermined probe region to form a hybrid satisfying the aforementioned conditions.

## EXAMPLES

The present invention will be described in detail below by way of Examples. Examples described below are preferred



embodiments according to the present invention and will not limit a technical scope of the present invention.

### Example 1

#### I. PCR for pUC118 EcoRI/BAP

##### (1) Primer Design

As a model sample having a target sequence, a commercially available vector, pUC118 EcoRI/BAP of 3162 bp (manufactured by Takara) was chosen. Three primers having the following sequences were designed for the vector. The base-sequence data of full-length pUC118 EcoRI/BAP was given by Takara and also available from a database known to the public.

Primers were designed in consideration of base sequence, GC % and melting temperature ( $T_m$ ) so as to specifically and efficiently amplify a desired portion of pUC118 EcoRI/BAP by PCR amplification.

Three primers: two forward primers (F1, F2) and one reverse primer (R1), were designed. When PCR amplification was performed by using pUC118 EcoRI/BAP as a template, and a set of primers, that is, a combination of F1 and R1 or F2 and R1, a PCR product of 1324 bp (PCR product 1) and a PCR product of 940 bp (PCR product 2) were obtained. The base sequence of each primer designed herein and the melting temperature ( $T_m$ ) of corresponding hybrid are shown in Table 1.

TABLE 1

Name	Base sequence	$T_m$ ( $^{\circ}$ C.)
F1	5' TGATTTGGGTGATGGTTCACGTAG 3' (SEQ ID NO:1)	60.9
F2	5' GAGCTGCATGTGTCAGAGGTTT 3' (SEQ ID NO:2)	61.0
R1	5' ATCAGCAATAAACCCAGCCAGCC 3' (SEQ ID NO:3)	61.5

The positional relationship between three primers and the full-length pUC118 EcoRI/BAP are shown in FIG. 2. The arrow of each primer is directed from the 5' to 3'.

##### (2) Synthesis of Primer

Three primers designed in Example 1 (1) were synthesized. Each primer was synthesized by a DNA synthesizer in accordance with a conventional method so as to have the designed base sequence. The DNA strands thus obtained were purified by cartridge purification to obtain three primers, each of which was diluted with a TE buffer to 10  $\mu$ M.

##### (3) PCR Amplification Reaction

A PCR amplification reaction was performed by using three primer synthesized in Example 1 (2), a vector, pUC118 EcoRI/BAP (manufactured by Takara), as a template DNA, and a PCR kit, HotStarTaq Master Mix (manufactured by QIAGEN). To the MasterMix containing 4 types of deoxy-nucleotides, dATP, dCTP, dTTP and dGTP, Cy3dUTP (manufactured by Amersham Biosciences) was added to fluorescent label the PCR product.

PCR amplification was performed by using two sets of a forward primer and a reverse primer combinations described in Example 1 (1). PCR was performed in accordance with the conditions shown in the protocol below. PCR solutions (1) and (2) were prepared in accordance with the recipe shown in Table 2 below.

TABLE 2

Composition of reaction solution	
Component	Content
Master Mix (manufactured by QIAGEN)	25 $\mu$ l
Template DNA (diluted pUC118, Manufactured by Takara)	1 $\mu$ l (10 ng)
Forward Primer F1 (Solution 1)	2.5 $\mu$ l (25 pmol/tube)
or	
Forward Primer F2 (solution 2)	
Reverse Primer(R1)	2.5 $\mu$ l (25 pmol/tube)
Cy3 dUTP(1 mM, manufactured by Amersham Biosciences)	2 $\mu$ l (25 pmol/tube)
H <sub>2</sub> O	17 $\mu$ l
Total	50 $\mu$ l

Each reaction solution prepared above was subjected to a PCR amplification reaction using a commercially available thermal cycler and in accordance with the temperature cycle protocol shown in Table 3 below. To explain more specifically, the reaction solution was held at 95 $^{\circ}$  C. for 15 minutes, subjected to denaturation at 92 $^{\circ}$  C. for 45 seconds, annealing at 55 $^{\circ}$  C. for 45 seconds, and extension at 72 $^{\circ}$  C. for 45 seconds. This cycle was repeated 25 times. Finally, the reaction solution was allowed to stand at 72 $^{\circ}$  C. for 10 minutes.

TABLE 3

Temperature conditions for PCR amplification reaction			
Step	Temperature	Holding time	Repetition
1	95 $^{\circ}$ C.	15 min.	
2	92 $^{\circ}$ C. (denaturation)	45 sec.	25 cycles
3	55 $^{\circ}$ C. (annealing)	45 sec.	
4	72 $^{\circ}$ C. (elongation)	1 min.	
4	72 $^{\circ}$ C.	10 min.	

After completion of the PCR, an amplified product was purified by use of a purification column (Qiagen QIAquick PCR Purification Kit). After the purification, the volume of a solution containing the PCR amplification product was adjusted to 50  $\mu$ l. The PCR amplification product solution was purified, an aliquot was taken therefrom, and then, electrophoretically analyzed in accordance with a conventional method. Presence of a band of desired base chain length was confirmed in each of the PCR products (1) and (2).

## II. Preparation of DNA Microarray

### (1) Design and Synthesis of Probe

Three types of probes were designed for the PCR product 1 obtained above. Each of the probes was designed in the same manner as with primer design so as to specifically recognize a certain base sequence (target sequence) in the sample (PCR product 1) and to have the similar GC % and hybrid melting temperature ( $T_m$ ).

The base sequences and  $T_m$  values of designed probes are shown in Table 4.

TABLE 4

Name	Base sequence	$T_m$ ( $^{\circ}$ C.)
P1	5' ATGGTGCACCTCTCAGTACAATCTGC 3' SEQ ID NO:4	75.7



TABLE 4-continued

Name	Base sequence	T <sub>m</sub> (° C.)
P2	5' GTGGGTTACATCGAACTGGATCTCA 3' SEQ ID NO:5	75.8
P3	5' GATAAAGTTGCAGGACCACTTCTGC 3' SEQ ID NO:6	75.7

In this probe design, the DNA strand that hybridizes with the probe was a strand extending from R1 primer.

Probes P2 and P3 can also form hybrids with the PCR product 2.

In the hybrids between PCR products 1 and 2 and the corresponding probes, the length of the 5' portion of the target strand upstream the hybrid region is represented by L1 and that of 3' portion downstream of the hybrid region is represented by L2. The values of L1, L2 and L1/L2 with respect to each of the PCR products are shown in Table 5.

TABLE 5

PCR product	Probe	L1	L2	L1/L2
1 F1-R1	P1	1053	246	4.28
	P2	545	754	0.72
	P3	35	1264	0.03
2 F2-R1	P2	545	370	1.47
	P3	35	880	0.04

The positional relationship between three types of probes, three primers and full-length pUC118 EcoRI/BAP is shown in FIG. 3.

### (2) Cleaning of Glass Substrate

A glass substrate made of synthetic quartz of 25 mm width×75 mm length×1 mm thickness (manufactured by Iiyama Tokushu Glass) was placed in a heat and alkali resistant rack and soaked in an ultrasonic washing solution prepared at a predetermined concentration. After the substrate was allowed to stand overnight in the washing solution, it was subjected to ultrasonic washing for 20 minutes. Thereafter, the glass substrate was taken out and lightly rinsed with pure water and then ultrasonically washed in ultrapure water for 20 minutes. The glass substrate was then soaked in 1N aqueous sodium hydroxide solution heated to 80° C. for 10 minutes and subjected again to pure water washing and ultrapure water washing steps. In this manner, a clean quartz glass substrate for use in a DNA microarray was prepared.

### (3) Surface Treatment

A silane coupling agent, KBM-603 (manufactured by Shin-Etsu Silicone) was dissolved in pure water to a concentration of 1%. The resultant solution was stirred at room temperature for 2 hours. Subsequently, the clean quartz substrate was soaked in the aqueous solution of the silane coupling agent and allowed to stand for 20 minutes at room temperature. The glass substrate was taken out, lightly washed with pure water, and dried by blowing with nitrogen gas to both surfaces of the substrate. The glass substrate dried with nitrogen gas was baked in an oven at 120° C. for one hour to complete the treatment with the coupling agent. In this manner, an amino group derived from silane coupling agent was introduced into the surfaces of the glass substrate.

While, N-(6-maleimidocaproyloxy)succinimide (hereinafter simply referred to as "EMCS") manufactured by Doujin Kagaku Kenkyusho was dissolved in a solvent mixture containing dimethylsulfoxide and ethanol in a ratio of 1:1 to a final concentration of 0.3 mg/ml to prepare an EMCS solution. After baking, the treated glass substrate was allowed to

cool and soaked in the EMCS solution prepared above at room temperature for 2 hours. During the soaking step, the amino group introduced into the glass substrate was allowed to react with a succinimide group derived from EMCS, thereby introducing a maleimide group derived from EMCS to the surfaces of the glass substrate. The glass substrate was taken out from the EMCS solution, washed with a solvent mixture of dimethylsulfoxide and ethanol mentioned above, washed further with ethanol and allowed to dry in a nitrogen gas atmosphere.

### (4) Synthesis of Probe DNA

Three types of probes designed in Step (1) above were synthesized.

To covalently bond a probe DNA to the glass substrate having a maleimide group introduced into the surface thereof, the 5' end of the probe DNA was modified with a thiol group in accordance with a conventional method. Thereafter, the protecting group that had been introduced to prevent side reactions during DNA synthesis was removed and HPLC purification and desalination treatment were performed.

The obtained probe DNA was dissolved in pure water and aliquots were subjected to lyophilization to remove water.

### (5) Ejection of Probe DNA by BJ Printer and Binding to Substrate Surface

An aqueous solution was prepared containing 7.5% by weight of glycerin, 7.5% by weight of thiodiglycol, 7.5% by weight of urea, and 1.0% by weight of acetylenol EH (manufactured by Kawaken Fine Chemicals Co., Ltd.). Subsequently, an aliquot of the probe DNA prepared above was dissolved in the solvent mixture prepared above to a final concentration of 10 μM. The resultant probe DNA solution was loaded in an ink tank of a bubble-jet printer, BJF-850 (trade name, manufactured by Canon Inc.), and the tank was installed in a print head.

Note that the bubble-jet printer was modified so as to print on a flat plate. The modified bubble-jet printer can spot liquid drops of a DNA solution (about 5 pl per drop) at about 120 μm pitch by inputting a printing pattern according to a predetermined file preparation method.

Using the modified bubble-jet printer, a probe DNA solution was spotted on a glass-substrate surface. A print pattern was previously designed so as to spot 16 spots for each probe on one DNA microarray and inkjet printing was performed. After spotting of the probe DNA solution onto a substrate in a desired pattern was confirmed by a magnifier, the microarray was left alone in a humidified chamber at normal temperature for 30 minutes to allow a maleimide group on the surface of a glass substrate to react with a thiol group at the 5' end of probe DNA.

### (6) Washing

After the reaction was performed for 30 minutes in the humidified chamber, probe DNA remaining unreacted on the glass-substrate surface was washed out by a 10 mM phosphate buffer (pH 7.0) containing 100 mM NaCl. In this way, a DNA microarray was obtained which had 16 spots of predetermined single-stranded probe DNA immobilized on the glass-substrate surface.

## III. Hybridization Reaction

Hybridization was performed by use of two types of PCR amplification products prepared as a sample nucleic acid in Step I on the DNA microarray prepared in Step II.

### (1) Blocking of DNA Microarray

Bovine serum albumin (BSA, fraction V, manufactured by Sigma) was dissolved in a 10 mM phosphate buffer containing 100 mM NaCl to make a concentration of 1% by weight.



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To this solution, the DNA microarray prepared in Step II was soaked at room temperature for 2 hours to block the glass-substrate surface. After completion of the blocking, the substrate was washed with 2×SSC solution [(300 mM of NaCl, 30 mM of sodium citrate (trisodium citrate dehydrate C<sub>6</sub>H<sub>5</sub>Na<sub>3</sub>·2H<sub>2</sub>O), pH 7.0)] containing 0.1% by weight sodium dodecyl sulfate (SDS) and then rinsed with pure water. Thereafter, the DNA microarray was dewatered by a spin dryer.

## (2) Preparation of Hybridization Solution

PCR products were prepared so as to have the same molar concentration. Hybridization solutions were prepared by using the PCR amplification product solutions so as to obtain final concentrations mentioned below.

## &lt;Hybridization Solution&gt;

6×SSPE/10% formamide/PCR amplification product solution (6×SSPE contains 900 mM NaCl, 60 mM NaH<sub>2</sub>PO<sub>4</sub>H<sub>2</sub>O, and 6 mM EDTA, pH 7.4).

## (3) Hybridization

The microarray dewatered was placed in a hybridization apparatus (Hybridization Station, manufactured by Genomic Solutions Inc.) and hybridization was performed using the hybridization solution of the aforementioned composition in accordance with the conditions and procedure described below.

## &lt;Conditions and Procedure of Hybridization&gt;

The hybridization solution was heated to 6.5° C. and allowed to stand for 3 minutes, subsequently at 92° C. for 2 minutes, and then at 55° C. for 4 hours. Thereafter, the DNA microarray was washed with 2×SSC, 0.1% SDS at 25° C., further with 2×SSC at 20° C., and if necessary, rinsed with pure water in accordance with a conventional manual, and finally dewatered by a spin drier to dry.

TABLE 6

Operation	Operation procedure/conditions
Reaction	65° C. 3 min → 92° C. 2 min → 55° C. 4 h
Wash	2 × SSC/0.1% SDS at 25° C. 2 × SSC at 20° C.
(Rinse)	H <sub>2</sub> O (manual rinsing)
Dry	Spin dry

## (4) Fluorescence Measurement

After completion of the hybridization reaction, the DNA microarray spin-dried was subjected to fluorescence measurement using a DNA-microarray fluorescence detection device (Genepix 4000B, manufactured by Axon Instruments) to measure fluorescence emitted from a hybrid. The fluorescent intensity values of PCR amplification products 1 and 2 measured with respect to individual probes are shown in Table 7 below.

In computation of fluorescent intensity, the fluorescent intensity of the portion where no DNA-probe spot was present was subtracted as a background value from the apparent fluorescent intensity of each DNA-probe spot. Measurement is performed twice and an average value thereof is shown.

TABLE 7

PCR product	Probe	Fluorescent intensity	L1/L2
1 F1-R1	P1	10	4.28
	P2	4607	0.72
	P3	7369	0.03

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TABLE 7-continued

PCR product	Probe	Fluorescent intensity	L1/L2
2 F2-R1	P2	725	1.47
	P3	1504	0.04

From the results above, with probes designed to have virtually the same T<sub>m</sub> value with respect to the same sample nucleic acid, the fluorescent intensity greatly differs with the probes. This fact suggests that the stability of hybrids greatly differs. High fluorescent value means that many hybrids are formed, in other words, hybrids are stably formed. The stability of a hybrid varies depending upon the ratio of L1/L2 (L1 is the length of 5' portion upstream the hybrid portion of the target, and L2 is the length of 3' portion downstream the hybridized portion of the target stand of the sample). It was found out that a hybrid was more stable when L1 is shorter than L2.

## Example 2

## I. PCR for pUC118 EcoRI/BAP

## (1) Primer Design

As a model sample having a target sequence, a commercially available vector, pUC118 EcoRI/BAP of 3162 bp (manufactured by Takara) was chosen. Primers having the following sequences were designed for the vector. The base-sequence data of full-length pUC118 EcoRI/BAP was given by Takara and also available from a database known to the public.

Primers were designed in consideration of base sequence, GC % and melting temperature (T<sub>m</sub>) so as to specifically and efficiently amplify desired portions of pUC118 EcoRI/BAP by PCR amplification.

Fifteen types of primers including 8 forward primers (F1, F2) and 7 reverse (R1) primers were designed. PCR amplification was performed using various combination of forward primers and reverse primers and pUC118 EcoRI/BAP as a template. In this manner, when PCR amplification was performed with respect to all combinations of 8 forward primers and 7 reverse primers, various PCR products having a length ranging from 104 bp to 2345 bp were produced.

The base sequences of primers thus designed are shown in Table 8.

TABLE 8

Name	Sequence (5' → 3')
FP1	CTTAATCGCCTTGCAGCACATC SEQ ID NO: 7
FP2	TGATTTGGGTGATGGTTCACGTAG SEQ ID NO: 8
FP3	GAGCTGCATGTGTCAGAGGTTT SEQ ID NO: 9
FP4	TTCTTAGACGTCAGGTGGCACT SEQ ID NO: 10
FP5	CCTTCCTGTTTTTGCTCACCCA SEQ ID NO: 11
FP6	GCATCTTACGGATGGCATGACA SEQ ID NO: 12

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TABLE 8-continued

Name	Sequence (5' → 3')
FP7	TGAAGCCATACCAAACGACGAG SEQ ID NO: 13
FP8	TTACTCTAGCTTCCCGCAACA SEQ ID NO: 14
RP1	TTCGGTGATGACGGTGAAAACC SEQ ID NO: 15
RP2	ACTGGTGAGTACTCAACCAAGTCA SEQ ID NO: 16
RP3	ATCAGCAATAAACCCAGCCAGCC SEQ ID NO: 17
RP4	TACGGGAGGGCTTACCATCTG SEQ ID NO: 18
RP5	CATCCATAGTTGCCTGACTCCC SEQ ID NO: 19
RP6	ACGCTCAGTGGAAACGAAAACCTC SEQ ID NO: 20
RP7	GCGCCTTATCCGGTAACTATCG SEQ ID NO: 21

Schematic positional-relationship between 15 types of primers and pUC118 EcoRI/BAP are shown in FIG. 4. The arrow on each primer indicates direction of 5' to 3'.

### (2) Synthesis of Primer

Fifteen primers designed in Example 2 (1) were synthesized. Each primer was synthesized by a DNA synthesizer in accordance with a conventional method so as to have the designed base sequence. The DNA strands thus obtained were purified by cartridge purification to obtain three primers, each of which was diluted with a TE buffer to 10 μM.

### (3) PCR Amplification Reaction

PCR amplification reaction was performed by using 15 types of primers synthesized in Example 2 (2), a vector, pUC118 EcoRI/BAP (manufactured by Takara), as a template DNA, and a PCR kit, AccuPrime Taq DNA Polymerase System (manufactured by Invitrogen). To a buffer in the AccuPrime kit containing 4 types of deoxynucleotides, dATP, dCTP, dTTP and dGTP, Cy3dUTP (manufactured by Amersham Biosciences) was added to fluorescent label the PCR product.

PCR amplification was performed with respect to 24 combinations of forward primers and reverse primers as shown in Table 9.

TABLE 9

No.	Forward primer	Reverse primer	Length of PCR product
1	FP1	RP1	811
2	FP1	RP2	1336
3	FP1	RP4	1786
4	FP1	RP6	2019
5	FP1	RP7	2345
6	FP2	RP4	1398
7	FP2	RP7	1957
8	FP3	RP4	1014
9	FP3	RP6	1247
10	FP3	RP7	1573
11	FP4	RP4	909
12	FP5	RP3	640
13	FP5	RP4	714
14	FP6	RP3	367

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TABLE 9-continued

No.	Forward primer	Reverse primer	Length of PCR product
15	FP6	RP4	441
16	FP6	RP7	1000
17	FP7	RP3	190
18	FP7	RP4	264
19	FP7	RP5	305
20	FP7	RP6	497
21	FP8	RP3	104
22	FP8	RP4	178
23	FP8	RP5	219
24	FP8	RP6	411

The reaction conditions are as shown in the protocol below. PCR solutions were prepared in accordance with the recipe shown in Table 10 below.

TABLE 10

Component	Content
AccuPrime Taq DNA Polymerase	0.5 μl
Template DNA (diluted pUC118, Manufactured by Takara) (100 pg/μl)	0.5 μl (50 pg)
Forward primer (10 μM)	1.25 μl
Reverse primer (10 μM)	1.25 μl
Cy3 dUTP (1 mM, manufactured by Amersham Biosciences)	0.5 μl
10 × AccuPrime PCR Buffer I	2.5 μl
H <sub>2</sub> O	18.5 μl
Total	25 μl

The reaction solution prepared above was subjected to a PCR amplification reaction using a commercially available thermal cycler in accordance with the temperature cycle protocol shown in Table 11. More specifically, the reaction solution was held at 94° C. for 2 minutes, subjected to denaturation at 92° C. for 30 seconds, annealing at 55° C. for 45 seconds, and extension at 72° C. for 60 seconds. This cycle was repeated 30 times. Finally, the reaction solution was allowed to stand at 72° C. for 10 minutes.

TABLE 11

Step	Temperature	Holding time	Number of repeats
1	94° C.	2 minutes	
2	92° C. Denaturation	30 seconds	30 cycles
3	55° C. Annealing	45 seconds	
4	72° C. Extension	1 minute	
4	72° C.	10 minutes	

After completion of the reaction, a PCR amplification product was purified by use of a purification column (Qiagen QIAquick PCR Purification Kit). After the purification, the volume of a solution containing the PCR amplification product was adjusted to 50 μl. The PCR amplification product solution was purified, an aliquot was taken therefrom, and then, electrophoretically analyzed in accordance with a conventional method. Presence of a band having a desired base chain length was confirmed in each of the 24 types of PCR products, and the production amount of each PCR product was determined.



## II. Preparation of DNA Microarray

## (1) Design of Probe

To detect 24 types of PCR products, probes were designed to 3 regions on pUC118 EcoRI/BAP. FIG. 5 shows probe regions on pUC118 EcoRI/BAP. Three probes were designed for three regions so as to be mutually different in chain length, that is, probes of 25-mers, 40-mers and 60-mers, were designed. The probes were designed in the same manner as in primers so as to specifically recognize a certain base sequence (target sequence) and to have the similar GC % and hybrid melting temperature ( $T_m$ ).

Table 12 shows the name, base sequence and  $T_m$  value of each of the 9 types of designed probes.

TABLE 12

Name	Sequence (5' → 3')	$T_m$ value
A25	ATG GTG CAC TCT CAG TAC AAT CTG G SEQ ID NO: 22	75.7
B25	GTG GGT TAC ATC GAA CTC GAT CTC A SEQ ID NO: 23	75.8
C25	GAT AAA GTT GCA GGA CCA CTT CTG C SEQ ID NO: 24	75.7
A40	ATG GTG CAC TCT CAG TAC AAT CTG CTC TGA TGC CGC ATA G SEQ ID NO: 25	80.7
B40	AGT TGG GTG CAC GAG TGG GTT ACA TCG AAC TGG ATC TCA A SEQ ID NO: 26	80.9
C40	TAG ACT GGA TGG AGG CGG ATA AAG TTG CAG GAC CAC TTC T SEQ ID NO: 27	80.8
A60	TTT TAT GGT GCA CTC TCA GTA CAA TCT GCT CTG ATG CCG CAT AGT TAA GCC AGC CCC GAC SEQ ID NO: 28	83.3
B60	GTG GGT TAC ATC GAA CTG GAT CTC AAC AGC GGT AAG ATC CTT GAG AGT TTT CGC CCC GAA SEQ ID NO: 29	83.3
C60	CTT CCC GGC AAC AAT TAA TAG ACT GGA TGG AGG CGG ATA AAG TTG CAG GAC CAC TTC TGC SEQ ID NO: 30	83.3

Each of the aforementioned probes has a sequence on the same strand of pUC118 EcoRI/BAP as that the forward primer. Therefore, the sequence that hybridizes with each of the probes to form a double strand is that of a complementary strand extending from a reverse primer.

In the hybrid between each of the PCR products and the corresponding probe, the length of a target strand upstream of the hybrid region (L1) and the length downstream of the hybrid region (L2), and L1/L2 are shown in Tables below. Table 13 shows 25-mer probes, A25, B25 and C25, Table 14 40-mer probes, and Table 15 60-mer probes.

TABLE 13

No.	Forward primer	Reverse primer	Probe	Length of PCR product	L1 (bp)	L2 (bp)	L1/L2
1	FP1	RP1	A25	811	152	634	0.24
2	FP1	RP2	A25	1336	677	634	1.07
3	FP1	RP2	B25	1336	169	1142	0.15
4	FP1	RP4	A25	1786	1127	634	1.78

TABLE 13-continued

No.	Forward primer	Reverse primer	Probe	Length of PCR product	L1 (bp)	L2 (bp)	L1/L2	
5	5	FP1	RP4	B25	1786	619	1142	0.54
	6	FP1	RP4	C25	1786	109	1652	0.07
	7	FP1	RP6	A25	2019	1360	634	2.15
	8	FP1	RP6	B25	2019	852	1142	0.75
	9	FP1	RP6	C25	2019	342	1652	0.21
	10	FP1	RP7	A25	2345	1686	634	2.66
	11	FP1	RP7	B25	2345	1178	1142	1.03
	12	FP1	RP7	C25	2345	668	1652	0.40
	13	FP2	RP4	A25	1398	1127	246	4.58
	14	FP2	RP4	B25	1398	619	754	0.82
	15	FP2	RP4	C25	1398	109	1264	0.09
	16	FP2	RP7	A25	1957	1686	246	6.85
	17	FP2	RP7	B25	1957	1178	754	1.56
	18	FP2	RP7	C25	1957	668	1264	0.53
	19	FP3	RP4	B25	1014	619	370	1.67
	20	FP3	RP4	C25	1014	109	880	0.12
	21	FP3	RP6	B25	1247	852	370	2.30
	22	FP3	RP6	C25	1247	342	880	0.39
	23	FP3	RP7	B25	1573	1178	370	3.18
	24	FP3	RP7	C25	1573	668	880	0.76
	25	FP4	RP4	B25	909	619	265	2.34
	26	FP4	RP4	C25	909	109	775	0.14
	27	FP5	RP3	B25	640	545	70	7.79
	28	FP5	RP3	C25	640	35	580	0.06
	29	FP5	RP4	B25	714	619	70	8.84
	30	FP5	RP4	C25	714	109	580	0.19
	31	FP6	RP3	C25	367	35	307	0.11
	32	FP6	RP4	C25	441	109	307	0.36
	33	FP6	RP7	C25	1000	668	307	2.18
	34	FP7	RP3	C25	190	35	130	0.27
	35	FP7	RP4	C25	264	109	130	0.84
	36	FP7	RP5	C25	305	150	130	1.15
	37	FP7	RP6	C25	497	342	130	2.63
	38	FP8	RP3	C25	104	35	44	0.80
	39	FP8	RP4	C25	178	109	44	2.48
	40	FP8	RP5	C25	219	150	44	3.41
	41	FP8	RP6	C25	411	342	44	7.77

TABLE 14

No.	Forward primer	Reverse primer	Probe	Length of PCR product	L1 (bp)	L2 (bp)	L1/L2
1	FP1	RP1	A40	811	137	634	0.22
2	FP1	RP2	A40	1336	662	634	1.04
3	FP1	RP2	B40	1336	168	1128	0.15
4	FP1	RP4	A40	1786	1112	634	1.75
5	FP1	RP4	B40	1786	618	1128	0.55
6	FP1	RP4	C40	1786	111	1635	0.07
7	FP1	RP6	A40	2019	1345	634	2.12
8	FP1	RP6	B40	2019	851	1128	0.75
9	FP1	RP6	C40	2019	344	1635	0.21
10	FP1	RP7	A40	2345	1671	634	2.64
11	FP1	RP7	B40	2345	1177	1128	1.04
12	FP1	RP7	C40	2345	670	1635	0.41
13	FP2	RP4	A40	1398	1112	246	4.52
14	FP2	RP4	B40	1398	618	740	0.84
15	FP2	RP4	C40	1398	111	1247	0.09
16	FP2	RP7	A40	1957	1671	246	6.79
17	FP2	RP7	B40	1957	1177	740	1.59
18	FP2	RP7	C40	1957	670	1247	0.54
19	FP3	RP4	B40	1014	618	356	1.74
20	FP3	RP4	C40	1014	111	863	0.13
21	FP3	RP6	B40	1247	851	356	2.39
22	FP3	RP6	C40	1247	344	863	0.40
23	FP3	RP7	B40	1573	1177	356	3.31
24	FP3	RP7	C40	1573	670	863	0.78
25	FP4	RP4	B40	909	618	251	2.46
26	FP4	RP4	C40	909	111	758	0.15
27	FP5	RP3	B40	640	545	70	7.79
28	FP5	RP3	C40	640	37	563	0.07
29	FP5	RP4	B40	714	618	56	11.04
30	FP5	RP4	C40	714	111	563	0.20
31	FP6	RP3	C40	367	37	290	0.13



TABLE 14-continued

No.	Forward primer	Reverse primer	Probe	Length of PCR product	Length of		L1/L2
					L1 (bp)	L2 (bp)	
32	FP6	RP4	C40	441	111	290	0.38
33	FP6	RP7	C40	1000	670	290	2.31
34	FP7	RP3	C40	190	37	113	0.33
35	FP7	RP4	C40	264	111	113	0.98
36	FP7	RP5	C40	305	152	113	1.35
37	FP7	RP6	C40	497	344	113	3.04
38	FP8	RP3	C40	104	37	27	1.37
39	FP8	RP4	C40	178	111	27	4.11
40	FP8	RP5	C40	219	152	27	5.63
41	FP8	RP6	C40	411	344	27	12.74

TABLE 15

No.	Forward primer	Reverse primer	Probe	Length of PCR product	Length of		L1/L2
					L1 (bp)	L2 (bp)	
1	FP1	RP1	A60	811	115	636	0.18
2	FP1	RP2	A60	1336	640	636	1.01
3	FP1	RP2	B60	1336	148	1128	0.13
4	FP1	RP4	A60	1786	1090	636	1.71
5	FP1	RP4	B60	1786	598	1128	0.53
6	FP1	RP4	C60	1786	91	1635	0.06
7	FP1	RP6	A60	2019	1323	636	2.08
8	FP1	RP6	B60	2019	831	1128	0.74
9	FP1	RP6	C60	2019	324	1635	0.20
10	FP1	RP7	A60	2345	1649	636	2.59
11	FP1	RP7	B60	2345	1157	1128	1.03
12	FP1	RP7	C60	2345	650	1635	0.40
13	FP2	RP4	A60	1398	1090	248	4.40
14	FP2	RP4	B60	1398	598	740	0.81
15	FP2	RP4	C60	1398	91	1247	0.07
16	FP2	RP7	A60	1957	1649	248	6.65
17	FP2	RP7	B60	1957	1157	740	1.56
18	FP2	RP7	C60	1957	650	1247	0.52
19	FP3	RP4	B60	1014	598	356	1.68
20	FP3	RP4	C60	1014	91	863	0.11
21	FP3	RP6	B60	1247	831	356	2.33
22	FP3	RP6	C60	1247	324	863	0.38
23	FP3	RP7	B60	1573	1157	356	3.25
24	FP3	RP7	C60	1573	650	863	0.75
25	FP4	RP4	B60	909	598	251	2.38
26	FP4	RP4	C60	909	91	758	0.12
27	FP5	RP3	B60	640	524	56	9.36
28	FP5	RP3	C60	640	17	563	0.03
29	FP5	RP4	B60	714	598	56	10.68
30	FP5	RP4	C60	714	91	563	0.16
31	FP6	RP3	C60	367	17	290	0.06
32	FP6	RP4	C60	441	91	290	0.31
33	FP6	RP7	C60	1000	650	290	2.24
34	FP7	RP3	C60	190	17	113	0.15
35	FP7	RP4	C60	264	91	113	0.81
36	FP7	RP5	C60	305	132	113	1.17
37	FP7	RP6	C60	497	324	113	2.87
38	FP8	RP3	C60	104	17	27	0.63
39	FP8	RP4	C60	178	91	27	3.37
40	FP8	RP5	C60	219	132	27	4.89
41	FP8	RP6	C60	411	324	27	12.00

## (2) Preparation of Microarray

The steps from probe synthesis to microarray preparation were performed in the same manner as in Example 1. In this Example, a microarray was prepared for each of 9 probes. Similarly to Example 1, 16 spots were fixed on a single microarray.

## III. Hybridization Reaction

Using the DNA microarray prepared in Step II and 24 PCR amplification products prepared in Step I, 243 kinds of hybridization were performed on the microarrays as shown in Tables 13 to 15.

## (1) Blocking of DNA Microarray

Blocking was performed by using BSA in the same manner as in Example 1.

## (2) Preparation of Hybridization Solution

PCR products were prepared so as to have the same molar concentration. Hybridization solutions were prepared by using the PCR amplification product solutions so as to obtain final concentrations mentioned below.

## &lt;Hybridization Solution&gt;

6×SSPE/10% formamide/0.05% SDS/PCR amplification product solution (6×SSPE contains 900 mM NaCl, 60 mM NaH<sub>2</sub>PO<sub>4</sub>·H<sub>2</sub>O, and 6 mM EDTA, pH 7.4).

## (3) Hybridization

The microarray dewatered was placed in a hybridization apparatus (Hybridization Station, manufactured by Genomic Solutions Inc.) and hybridization was performed using the hybridization solution of the aforementioned composition in accordance with the conditions and procedure described below.

## &lt;Conditions and Procedure of Hybridization&gt;

The hybridization solution was added to a microarray at 65° C., allowed to stand at 92° C. for 2 minutes, and then at 55° C. for 4 hours. Thereafter, the microarray was washed with 2×SSC and 0.1% SDS at 25° C., further washed with 2×SSC at 20° C., optionally rinsed with pure water in accordance with a conventional manual, and finally dewatered by a spin drier to dry.

TABLE 16

Operation	Operation procedure/conditions
Addition	Hybridization solution is added at 65° C.
Reaction	92° C. 2 minutes → 55° C. 4 hours
Wash	2 × SSC/0.1% SDS at 25° C.
(Rinse)	2 × SSC at 20° C.
Dry	H <sub>2</sub> O(manual rinse) Spin dry

## (4) Fluorescence Measurement

After completion of the hybridization reaction, fluorescence was measured in the same manner as in Example 1. The fluorescent intensity values of individual PCR products are shown in Tables 17 to 19, which correspond to 25 mer, 40 mer, and 60 mer probes, respectively. Fluorescent intensity was calculated in the same manner as in Example 1.

TABLE 17

No.	Forward primer	Reverse primer	Probe	Length of PCR product	Length		L1/L2	Fluorescent intensity
					L1 (bp)	L2 (bp)		
1	FP1	RP1	A25	811	152	634	0.24	3172.0
2	FP1	RP2	A25	1336	677	634	1.07	52.4
3	FP1	RP2	B25	1336	169	1142	0.15	3638.8
4	FP1	RP4	A25	1786	1127	634	1.78	2.2
5	FP1	RP4	B25	1786	619	1142	0.54	3077.4
6	FP1	RP4	C25	1786	109	1652	0.07	3519.9
7	FP1	RP6	A25	2019	1360	634	2.15	0.6
8	FP1	RP6	B25	2019	852	1142	0.75	3020.5
9	FP1	RP6	C25	2019	342	1652	0.21	3449.8
10	FP1	RP7	A25	2345	1686	634	2.66	1.6
11	FP1	RP7	B25	2345	1178	1142	1.03	722.5
12	FP1	RP7	C25	2345	668	1652	0.40	2252.7
13	FP2	RP4	A25	1398	1127	246	4.58	1.5
14	FP2	RP4	B25	1398	619	754	0.82	2202.0
15	FP2	RP4	C25	1398	109	1264	0.09	2762.2
16	FP2	RP7	A25	1957	1686	246	6.85	1.4
17	FP2	RP7	B25	1957	1178	754	1.56	24.1
18	FP2	RP7	C25	1957	668	1264	0.53	1951.8
19	FP3	RP4	B25	1014	619	370	1.67	22.4



TABLE 17-continued

No.	Forward primer	Reverse primer	Probe	Length of PCR product	L1 (bp)	L2 (bp)	L1/L2	Fluorescent intensity
20	FP3	RP4	C25	1014	109	880	0.12	2763.2
21	FP3	RP6	B25	1247	852	370	2.30	0.5
22	FP3	RP6	C25	1247	342	880	0.39	2203.1
23	FP3	RP7	B25	1573	1178	370	3.18	0.5
24	FP3	RP7	C25	1573	668	880	0.76	519.7
25	FP4	RP4	B25	909	619	265	2.34	1.8
26	FP4	RP4	C25	909	109	775	0.14	1893.2
27	FP5	RP3	B25	640	545	70	7.79	1.1
28	FP5	RP3	C25	640	35	580	0.06	2009.7
29	FP5	RP4	B25	714	619	70	8.84	1.6
30	FP5	RP4	C25	714	109	580	0.19	1388.6
31	FP6	RP3	C25	367	35	307	0.11	1338.6
32	FP6	RP4	C25	441	109	307	0.36	484.5
33	FP6	RP7	C25	1000	668	307	2.18	1.3
34	FP7	RP3	C25	190	35	130	0.27	590.1
35	FP7	RP4	C25	264	109	130	0.84	73.4
36	FP7	RP5	C25	305	150	130	1.15	167.3
37	FP7	RP6	C25	497	342	130	2.63	2.7
38	FP8	RP3	C25	104	35	44	0.80	797.2
39	FP8	RP4	C25	178	109	44	2.48	709.4
40	FP8	RP5	C25	219	150	44	3.41	643.4
41	FP8	RP6	C25	411	342	44	7.77	3.1

TABLE 18

No.	Forward primer	Reverse primer	Probe	Length of PCR product	L1 (bp)	L2 (bp)	L1/L2	Fluorescent intensity
1	FP1	RP1	A40	811	137	634	0.22	6310.5
2	FP1	RP2	A40	1336	662	634	1.04	466.2
3	FP1	RP2	B40	1336	168	1128	0.15	7415.4
4	FP1	RP4	A40	1786	1112	634	1.75	10.0
5	FP1	RP4	B40	1786	618	1128	0.55	6451.3
6	FP1	RP4	C40	1786	111	1635	0.07	8547.4
7	FP1	RP6	A40	2019	1345	634	2.12	4.8
8	FP1	RP6	B40	2019	851	1128	0.75	5920.8
9	FP1	RP6	C40	2019	344	1635	0.21	8037.6
10	FP1	RP7	A40	2345	1671	634	2.64	3.0
11	FP1	RP7	B40	2345	1177	1128	1.04	1942.1
12	FP1	RP7	C40	2345	670	1635	0.41	6506.2
13	FP2	RP4	A40	1398	1112	246	4.52	3.2
14	FP2	RP4	B40	1398	618	740	0.84	5026.4
15	FP2	RP4	C40	1398	111	1247	0.09	7090.1
16	FP2	RP7	A40	1957	1671	246	6.79	3.7
17	FP2	RP7	B40	1957	1177	740	1.59	121.1
18	FP2	RP7	C40	1957	670	1247	0.54	5571.0
19	FP3	RP4	B40	1014	618	356	1.74	105.6
20	FP3	RP4	C40	1014	111	863	0.13	7291.5
21	FP3	RP6	B40	1247	851	356	2.39	4.0
22	FP3	RP6	C40	1247	344	863	0.40	6122.6
23	FP3	RP7	B40	1573	1177	356	3.31	0.5
24	FP3	RP7	C40	1573	670	863	0.78	1967.1
25	FP4	RP4	B40	909	618	251	2.46	19.9
26	FP4	RP4	C40	909	111	758	0.15	5674.5
27	FP5	RP3	B40	640	545	70	7.79	3.5
28	FP5	RP3	C40	640	37	563	0.07	4904.3
29	FP5	RP4	B40	714	618	56	11.04	4.0
30	FP5	RP4	C40	714	111	563	0.20	4248.7
31	FP6	RP3	C40	367	37	290	0.13	3611.1
32	FP6	RP4	C40	441	111	290	0.38	1810.2
33	FP6	RP7	C40	1000	670	290	2.31	7.4
34	FP7	RP3	C40	190	37	113	0.33	1390.3
35	FP7	RP4	C40	264	111	113	0.98	334.4
36	FP7	RP5	C40	305	152	113	1.35	579.7
37	FP7	RP6	C40	497	344	113	3.04	13.7
38	FP8	RP3	C40	104	37	27	1.37	1434.0
39	FP8	RP4	C40	178	111	27	4.11	1674.2
40	FP8	RP5	C40	219	152	27	5.63	1737.6
41	FP8	RP6	C40	411	344	27	12.74	14.6

TABLE 19

No.	Forward primer	Reverse primer	Probe	Length of PCR product	L1 (bp)	L2 (bp)	L1/L2	Fluorescent intensity
1	FP1	RP1	A60	811	115	636	0.18	9191.0
2	FP1	RP2	A60	1336	640	636	1.01	1963.3
3	FP1	RP2	B60	1336	148	1128	0.13	10343.4
4	FP1	RP4	A60	1786	1090	636	1.71	42.7
5	FP1	RP4	B60	1786	598	1128	0.53	11856.1
6	FP1	RP4	C60	1786	91	1635	0.06	10930.5
7	FP1	RP6	A60	2019	1323	636	2.08	10.7
8	FP1	RP6	B60	2019	831	1128	0.74	11330.2
9	FP1	RP6	C60	2019	324	1635	0.20	11745.8
10	FP1	RP7	A60	2345	1649	636	2.59	5.3
11	FP1	RP7	B60	2345	1157	1128	1.03	5436.4
12	FP1	RP7	C60	2345	650	1635	0.40	10108.0
13	FP2	RP4	A60	1398	1090	248	4.40	10.9
14	FP2	RP4	B60	1398	598	740	0.81	9617.4
15	FP2	RP4	C60	1398	91	1247	0.07	9193.6
16	FP2	RP7	A60	1957	1649	248	6.65	7.6
17	FP2	RP7	B60	1957	1157	740	1.56	826.5
18	FP2	RP7	C60	1957	650	1247	0.52	8783.0
19	FP3	RP4	B60	1014	598	356	1.68	1061.5
20	FP3	RP4	C60	1014	91	863	0.11	9409.1
21	FP3	RP6	B60	1247	831	356	2.33	44.6
22	FP3	RP6	C60	1247	324	863	0.38	9139.9
23	FP3	RP7	B60	1573	1157	356	3.25	0.8
24	FP3	RP7	C60	1573	650	863	0.75	3470.7
25	FP4	RP4	B60	909	598	251	2.38	306.8
26	FP4	RP4	C60	909	91	758	0.12	7428.8
27	FP5	RP3	B60	640	524	56	9.36	9.1
28	FP5	RP3	C60	640	17	563	0.03	5929.9
29	FP5	RP4	B60	714	598	56	10.68	11.9
30	FP5	RP4	C60	714	91	563	0.16	6037.3
31	FP6	RP3	C60	367	17	290	0.06	5069.8
32	FP6	RP4	C60	441	91	290	0.31	2892.6
33	FP6	RP7	C60	1000	650	290	2.24	15.4
34	FP7	RP3	C60	190	17	113	0.15	2119.3
35	FP7	RP4	C60	264	91	113	0.81	649.6
36	FP7	RP5	C60	305	132	113	1.17	1108.0
37	FP7	RP6	C60	497	324	113	2.87	37.4
38	FP8	RP3	C60	104	17	27	0.63	1448.2
39	FP8	RP4	C60	178	91	27	3.37	2060.3
40	FP8	RP5	C60	219	132	27	4.89	2130.9
41	FP8	RP6	C60	411	324	27	12.00	59.8

## IV. Analysis of Results

Based on the results, graphs were made by plotting fluorescent intensity to L1/L2 value. To facilitate understanding, the vertical and horizontal axes are graduated logarithmically. FIGS. 6, 7 and 8 show the cases of 25-mer, 40-mer and 60-mer probes, respectively.

From the results above, even when the same sample in the same concentration was reacted with the probes designed to have the same length and virtually the same T<sub>m</sub> value, the fluorescent intensities of the probes greatly differ. This means the stability of hybrids greatly differs. It was found out that if the L1/L2 value falls within the range of 0 to 1.5, a high fluorescent value was obtained, and thus a stable hybrid was obtained. If the L1/L2 value ranges from 0 to 1, the hybrid is particularly stable. In contrast, if the L1/L2 value is 2 or more, fluorescent intensity was extremely low with several exceptions, and the resultant hybrid had an extremely low stability.

## Example 3

## (1) Design of Probe

Of nine probes designed in Example 2, three types of 25-mer probes were modified to have a thiol group at the 3' end. A thiol group was introduced only to the 3' end and not to the 5' end.



Table 20 shows the name, base sequence, and T<sub>m</sub> value of each of the three types of probes.

TABLE 20

Name	Sequence (5' → 3')	T <sub>m</sub> value
AR25	ATGGTGCACACTGTGAGTAGAATGTGC SEQ ID NO: 24	75.7
BR25	GTGGGTTAGATGGAAGTGGATGTGA SEQ ID NO: 25	75.8
CR25	GATAAAGTTGCAGGAGCAGTTbTGG SEQ ID NO: 26	75.7

The same as in Example 2, the sequence that hybridizes with each of the probes to form a hybrid is the complementary strand extending from a reverse primer.

An example of hybrids formed between each of the PCR products (target strand) synthesized in Example 2 and each of the three probes synthesized herein, is shown in FIG. 9, where the length of a target strand upstream the hybridized region is represented by B<sub>5</sub> (L1) and that downstream the hybridized region is represented by B<sub>3</sub>(L2). The values of L1, L2 and L1/L2 with respect to each of the PCR products synthesized in Example 2 are shown in Table 21. Since a probe and each of PCR products (target strand) have the same positional relationship, the values of L1, L2 and L1/L2 are the same as in Example 2.

TABLE 21

No.	Forward primer	Reverse primer	Probe	Length of PCR product	L1 (bp)	L2 (bp)	L1/L2
1	FP1	RP1	AR25	811	152	634	0.24
2	FP1	RP2	AR25	1336	677	634	1.07
3	FP1	RP2	BR25	1336	169	1142	0.15
4	FP1	RP4	AR25	1786	1127	634	1.78
5	FP1	RP4	BR25	1786	619	1142	0.54
6	FP1	RP4	CR25	1786	109	1652	0.07
7	FP1	RP6	AR25	2019	1360	634	2.15
8	FP1	RP6	BR25	2019	852	1142	0.75
9	FP1	RP6	CR25	2019	342	1652	0.21
10	FP1	RP7	AR25	2345	1686	634	2.66
11	FP1	RP7	BR25	2345	1178	1142	1.03
12	FP1	RP7	CR25	2345	668	1652	0.40
13	FP2	RP4	AR25	1398	1127	246	4.58
14	FP2	RP4	BR25	1398	619	754	0.82
15	FP2	RP4	CR25	1398	109	1264	0.09
16	FP2	RP7	AR25	1957	1686	246	6.85
17	FP2	RP7	BR25	1957	1178	754	1.56
18	FP2	RP7	CR25	1957	668	1264	0.53
19	FP3	RP4	BR25	1014	619	370	1.67
20	FP3	RP4	CR25	1014	109	880	0.12
21	FP3	RP6	BR25	1247	852	370	2.30
22	FP3	RP6	CR25	1247	342	880	0.39
23	FP3	RP7	BR25	1573	1178	370	3.18
24	FP3	RP7	CR25	1573	668	880	0.76
25	FP4	RP4	BR25	909	619	265	2.34
26	FP4	RP4	CR25	909	109	775	0.14
27	FP5	RP3	BR25	640	545	70	7.79
28	FP5	RP3	CR25	640	35	580	0.06
29	FP5	RP4	BR25	714	619	70	8.84
30	FP5	RP4	CR25	714	109	580	0.19
31	FP6	RP3	CR25	367	35	307	0.11
32	FP6	RP4	CR25	441	109	307	0.36
33	FP6	RP7	CR25	1000	668	307	2.18
34	FP7	RP3	CR25	190	35	130	0.27
35	FP7	RP4	CR25	264	109	130	0.84
36	FP7	RP5	CR25	305	150	130	1.15
37	FP7	RP6	CR25	497	342	130	2.63
38	FP8	RP3	CR25	104	35	44	0.80
39	FP8	RP4	CR25	178	109	44	2.48
40	FP8	RP5	CR25	219	150	44	3.41
41	FP8	RP6	CR25	411	342	44	7.77

TABLE 21-continued

No.	Forward primer	Reverse primer	Probe	Length of PCR product	L1 (bp)	L2 (bp)	L1/L2
40	FP8	RP5	CR25	219	150	44	3.41
41	FP8	RP6	CR25	411	342	44	7.77

## (2) Preparation of Microarray

A DNA microarray was prepared for each of three probes in the same manner as in Example 2 and 16 spots were fixed on a single DNA microarray.

## (3) Hybridization Reaction and Measurement of Fluorescence

Hybridization reaction was performed and then fluorescence was measured in the same manner as in Example 2. The fluorescent intensity values obtained from individual microarrays are shown in Table 22.

TABLE 22

No.	Forward primer	Reverse primer	Probe	Length of PCR product	L1 (bp)	L2 (bp)	L1/L2	Fluorescent intensity
1	FP1	RP1	AR25	811	152	634	0.24	2569.3
2	FP1	RP2	AR25	1336	677	634	1.07	54.0
3	FP1	RP2	BR25	1336	169	1142	0.15	3056.6
4	FP1	RP4	AR25	1786	1127	634	1.78	1.8
5	FP1	RP4	BR25	1786	619	1142	0.54	2277.3
6	FP1	RP4	CR25	1786	109	1652	0.07	3379.1
7	FP1	RP6	AR25	2019	1360	634	2.15	0.5
8	FP1	RP6	BR25	2019	852	1142	0.75	2476.8
9	FP1	RP6	CR25	2019	342	1652	0.21	2207.9
10	FP1	RP7	AR25	2345	1686	634	2.66	1.4
11	FP1	RP7	BR25	2345	1178	1142	1.03	556.3
12	FP1	RP7	CR25	2345	668	1652	0.40	1667.0
13	FP2	RP4	AR25	1398	1127	246	4.58	1.3
14	FP2	RP4	BR25	1398	619	754	0.82	1431.3
15	FP2	RP4	CR25	1398	109	1264	0.09	2679.3
16	FP2	RP7	AR25	1957	1686	246	6.85	1.2
17	FP2	RP7	BR25	1957	1178	754	1.56	49.8
18	FP2	RP7	CR25	1957	668	1264	0.53	1678.5
19	FP3	RP4	BR25	1014	619	370	1.67	18.8
20	FP3	RP4	CR25	1014	109	880	0.12	2321.1
21	FP3	RP6	BR25	1247	852	370	2.30	0.0
22	FP3	RP6	CR25	1247	342	880	0.39	2137.0
23	FP3	RP7	BR25	1573	1178	370	3.18	0.4
24	FP3	RP7	CR25	1573	668	880	0.76	556.0
25	FP4	RP4	BR25	909	619	265	2.34	3.7
26	FP4	RP4	CR25	909	109	775	0.14	1666.0
27	FP5	RP3	BR25	640	545	70	7.79	0.8
28	FP5	RP3	CR25	640	35	580	0.06	1748.5
29	FP5	RP4	BR25	714	619	70	8.84	0.1
30	FP5	RP4	CR25	714	109	580	0.19	1222.0
31	FP6	RP3	CR25	367	35	307	0.11	1030.7
32	FP6	RP4	CR25	441	109	307	0.36	382.8
33	FP6	RP7	CR25	1000	668	307	2.18	1.1
34	FP7	RP3	CR25	190	35	130	0.27	454.4
35	FP7	RP4	CR25	264	109	130	0.84	67.6
36	FP7	RP5	CR25	305	150	130	1.15	140.6
37	FP7	RP6	CR25	497	342	130	2.63	2.9
38	FP8	RP3	CR25	104	35	44	0.80	693.5
39	FP8	RP4	CR25	178	109	44	2.48	524.9
40	FP8	RP5	CR25	219	150	44	3.41	540.4
41	FP8	RP6	CR25	411	342	44	7.77	4.8

## (4) Analysis of Results

A graph was made by plotting fluorescent intensity to L1/L2 value in the same manner as in Example 2 and shown



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in FIG. 10. In this Example also the vertical and horizontal axes are graduated logarithmically. As is apparent from this graph, high fluorescent values are obtained at an L1/L2 value from 0 to 1.5. It was clearly demonstrated that stable hybrids were obtained. The high intensity is remarkably obtained especially at an L1/L2 value from 0 to 1.

The results demonstrate that a stable hybrid is also formed, depending upon the relationship between L1 and L2, even when a probe immobilized at the 3' end through a thiol group, the same as the case immobilized at the 5' end. Hence, no

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matter what type of immobilization method is employed, a stable hybrid can be obtained according to the present invention.

The present invention is not limited to the above embodiments and various changes and modifications can be made within the spirit and scope of the present invention. Therefore to apprise the public of the scope of the present invention, the following claims are made.

This application claims priority from Japanese Patent Application No. 2003-324647 filed on Sep. 17, 2003, which is hereby incorporated by reference herein.

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What is claimed is:

1. A method for designing a nucleic acid probe set comprising more than one probe nucleic acid immobilized on a substrate for hybridizing with target sequences in target strands of sample nucleic acids, said method comprising:

confirming that 70% or more of the probes on the substrate satisfy the following conditions when the probes hybridize to the sample nucleic acids:

the probe nucleic acid is shorter than the target strand; the sample nucleic acid and a complementary probe sequence are present in the hybrid; and

$L1 \leq L2$ , wherein  $L1$  is the number of bases of a 5' portion of the target strand not being the target sequence,  $L2$  is the number of bases of a 3' portion of the target strand not being the target sequence, and each of  $L1$  and  $L2$  is greater than 0.

2. A method for designing a nucleic acid probe set comprising more than one probe nucleic acid immobilized on a

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substrate for hybridizing with target sequences in target strands of sample nucleic acids, said method comprising:

confirming that 70% or more of the probes on the substrate satisfy the following conditions (1) and (2):

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(1) the hybrid is formed between a target sequence present in a target strand of the sample nucleic acid and a complementary probe sequence present in the probe set; and

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(2) the target strand of the sample nucleic acid is longer than the probe nucleic acid and  $L1 \leq L2$ , wherein  $L1$  is the number of bases of a 5' portion of the target strand not being the target sequence,  $L2$  is the number of bases of a 3' portion of the target strand not being the target sequence, and each of  $L1$  and  $L2$  is greater than 0.

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